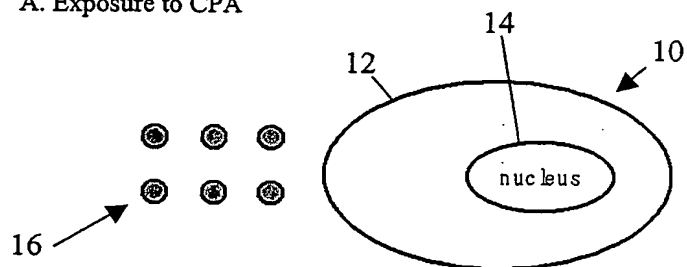
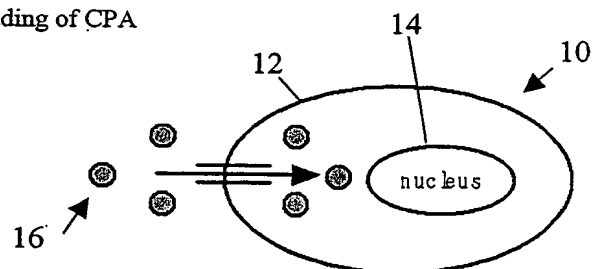


Sheet 1 of 9

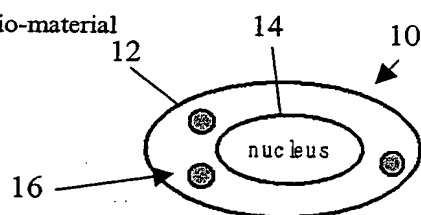
A. Exposure to CPA



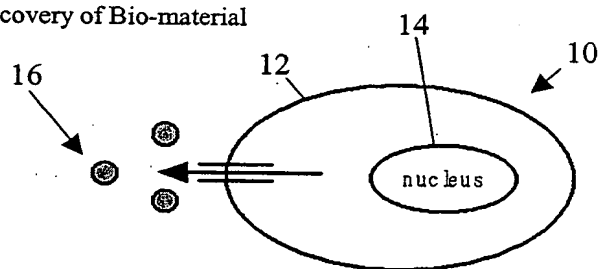
B. Loading of CPA



C. Preservation of Bio-material

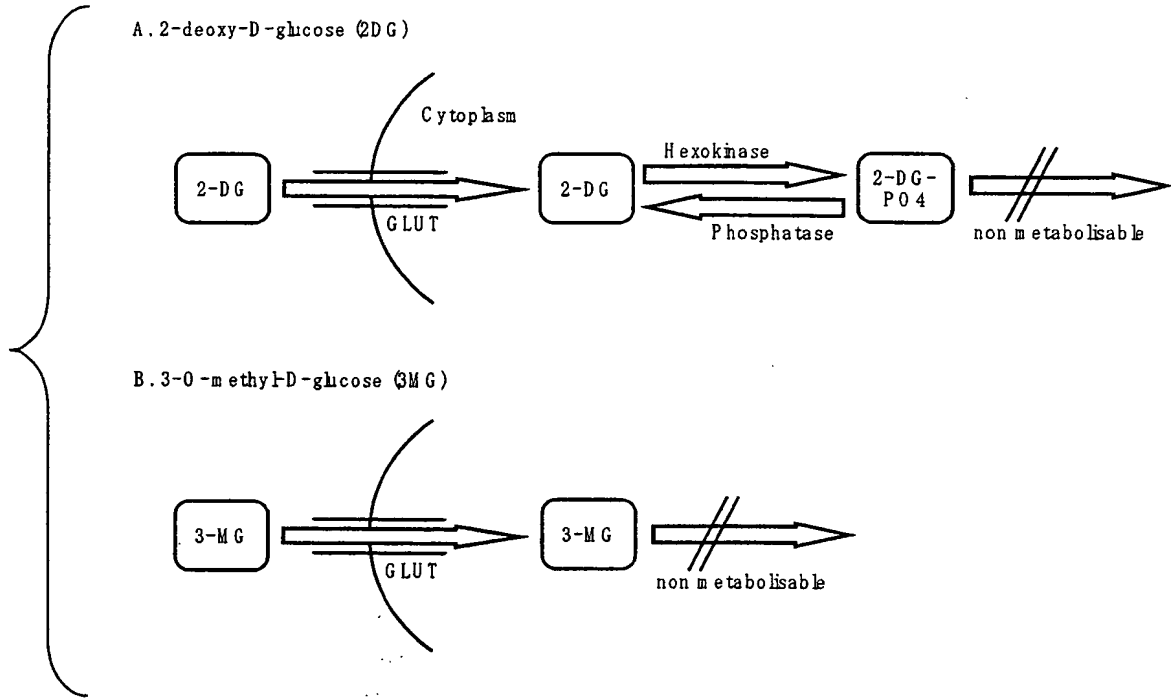


D. Recovery of Bio-material



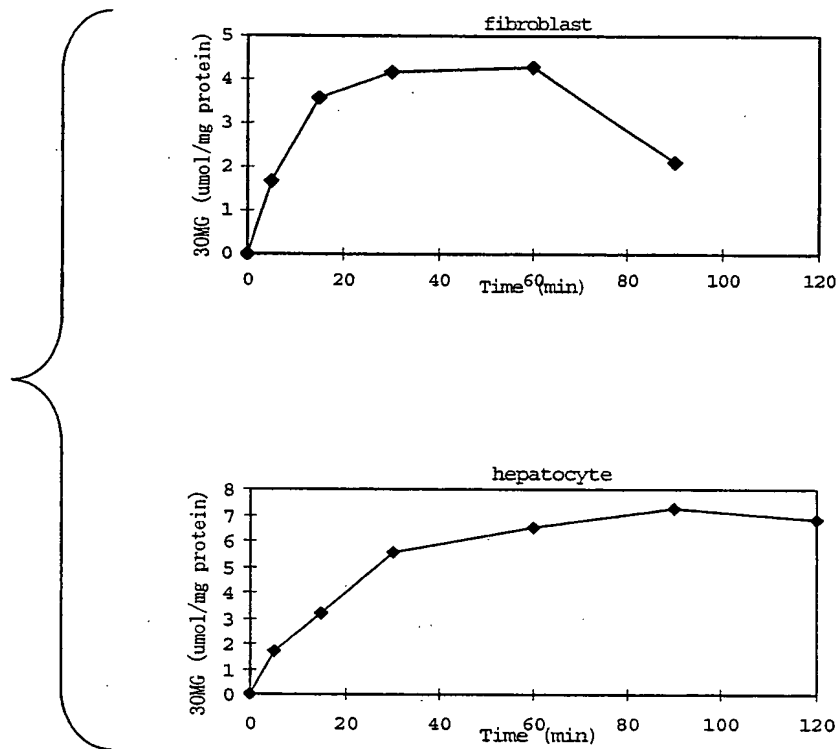
**FIGURE 1**

Sheet 2 of 9



**FIGURE 2**

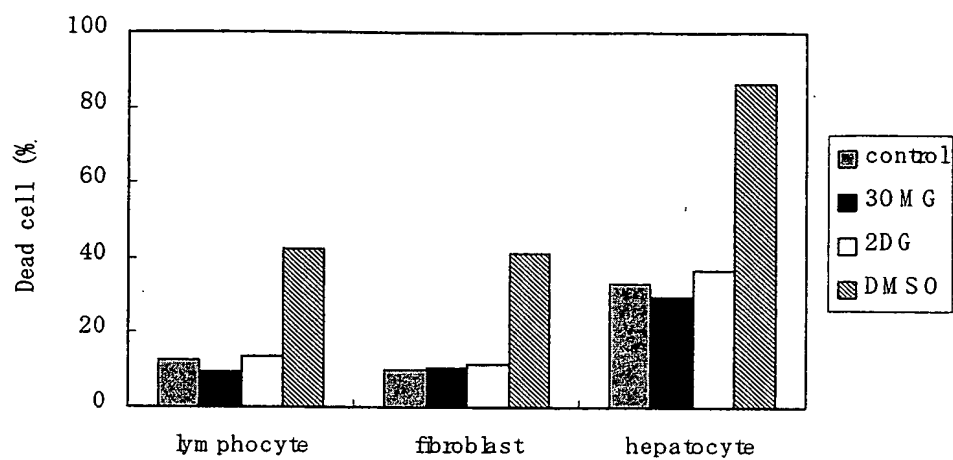
Sheet 3 of 9



Concentration of intracellular 3OMG measured using radiolabeled 3OMG

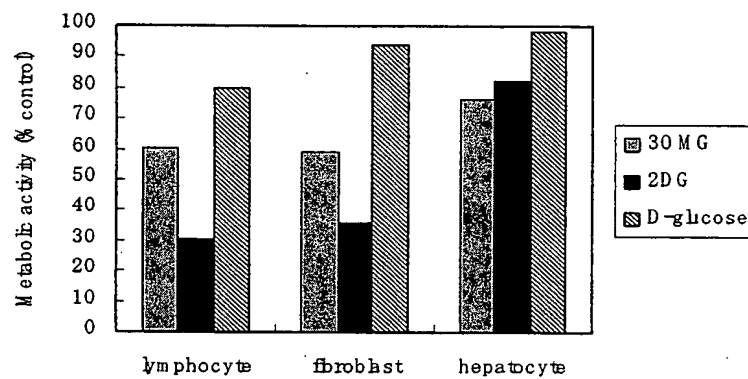
**FIGURE 3**

Sheet 4 of 9



Percentage of dead cells after loading glucoses or DMSO

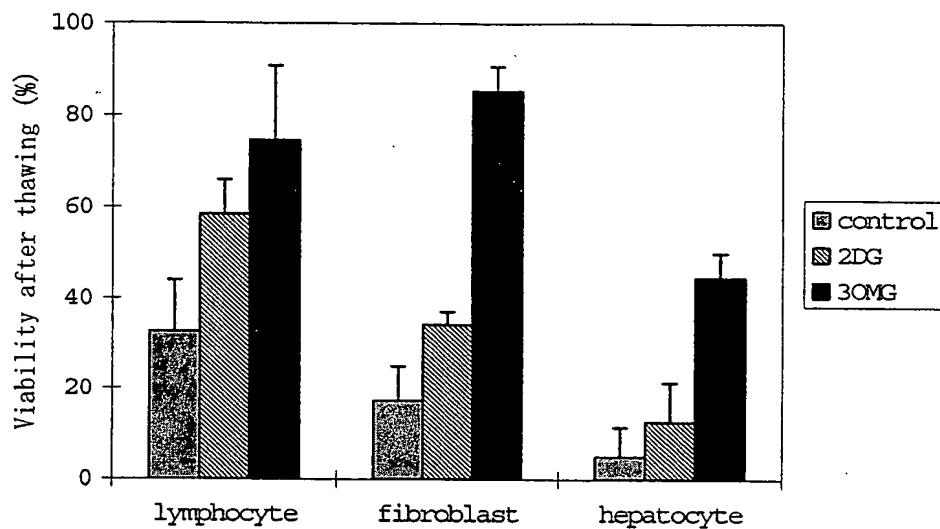
**FIGURE 4**



Metabolic activity was assessed measuring the MTT reduction activity

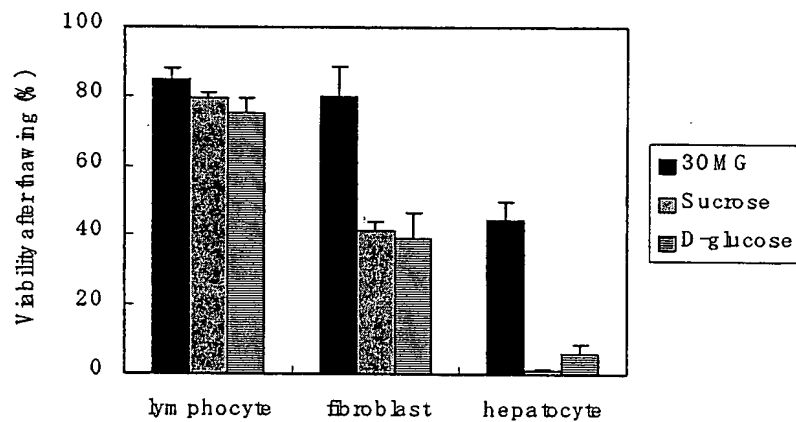
**FIGURE 5**

Sheet 5 of 9



Viability of cryopreserved mammalian cells with glucose loading

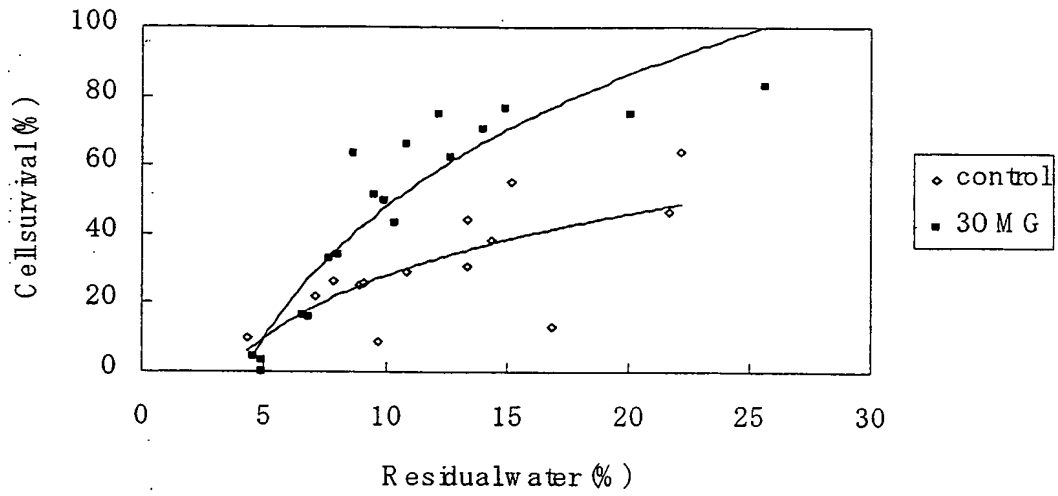
FIGURE 6



Viability of cryopreserved mammalian cells preserved using different agents

FIGURE 7

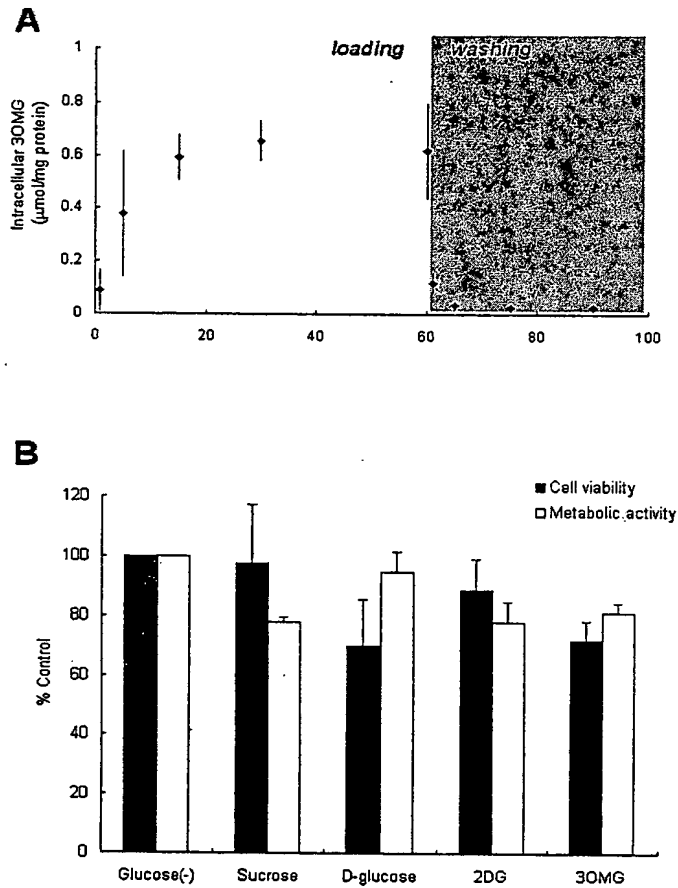
Sheet 6 of 9



Cell survival as a function of residual water in the sample after drying with or without 30MG loading

**FIGURE 8**

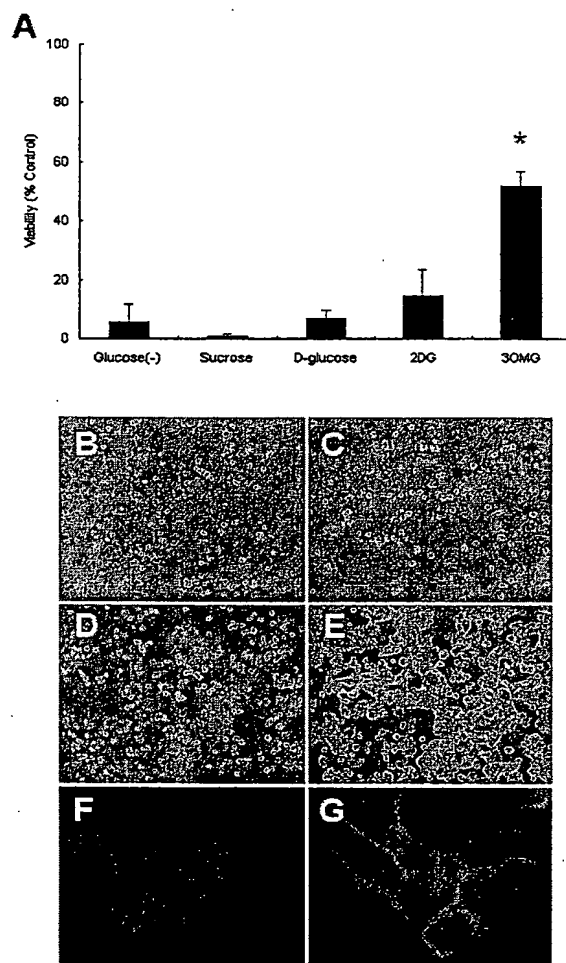
Sheet 7 of 9



Kinetics of 3OMG uptake and efflux on hepatocytes and effects on viability and metabolic activity

**FIGURE 9**

Sheet 8 of 9

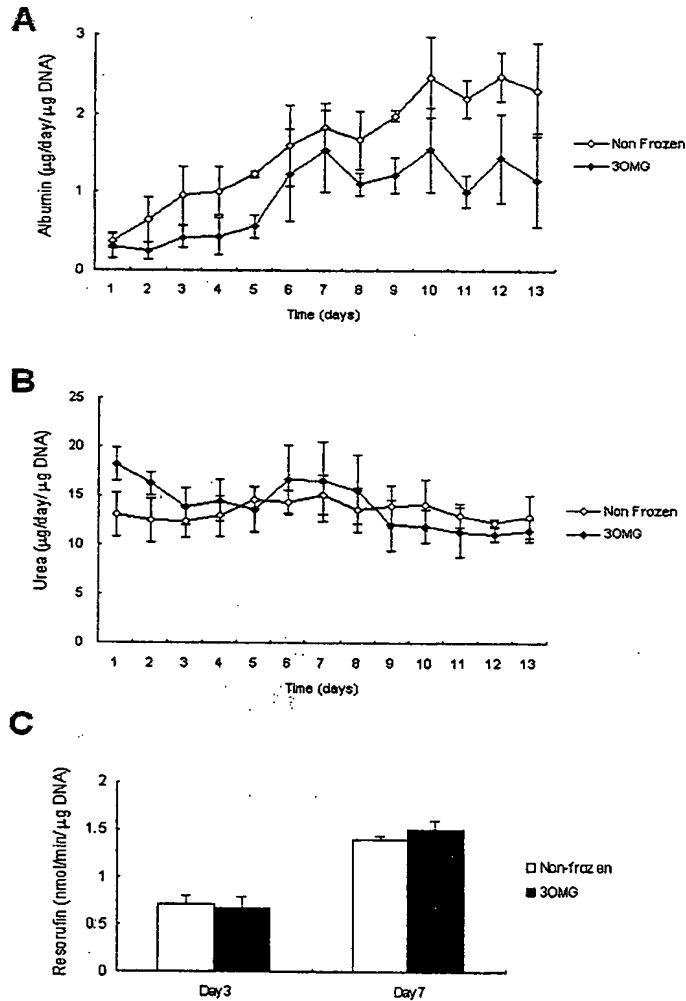


Post-thaw viability of cryopreserved hepatocytes

**FIGURE 10**



Sheet 9 of 9



Albumin production, urea production, and cytochrome P450 activity of treated hepatocytes

**FIGURE 11**